Hair zinc level in Down syndrome

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Abstract – Immunological, endocrinological, and haematological abnormalities are relatively common in people with Down syndrome (Cuadrado & Barrena, 1996; Decoq & Vincker, 1995; Hestnes et al., 1991; Sustrova & Strbak, 1994; Kempski, Chessells & Reeves, 1997; Kivivuori, Rajantie, & Siimes, 1996; David et al., 1996; Gjertson, Sturm & Berger, 1999). Zinc is one of the elements that act in the maintenance of normal function of these systems. This study was designed to investigate zinc levels in children with Down syndrome. Zinc levels were measured in hair using atomic absorption spectrophotometry. The hair zinc level of 19 children with Down syndrome was compared with the zinc level of 11 typically developing children. Hair zinc levels were found to be significantly lower (p < .05) in those with Down syndrome (average 95.18 ± 56.10 ppm) than in the typically developing children (average 208.88 ± 152.37 ppm). Some of the problems experienced by children with Down syndrome may be due to these low zinc levels, but further research is required to confirm these results, and to establish any correlation with these problems.

Keywords: Down syndrome, hair analysis, zinc

Introduction

Zinc is an important element in protein synthesis and gene expression involving the immune and endocrinological systems. Zinc deficiency may play a crucial role in some of the pathological manifestations associated with Down syndrome such as infections and malfunctioning of the thyroid gland (Bjorksten et al., 1980; Sustrova, & Strbak, 1994).

There have been a number of studies showing that low zinc levels occur in many children with Down syndrome and that oral zinc supplementation may be useful in correcting some of the immune and endocrinological disorders associated with thyroid dysregulation in these children (Franceschi et al., 1988; Kadrabova, Madaric, Sustrova & Ginter, 1996; Licastro et al., 1994; Licastro et al., 1992).

It has been reported that oral zinc supplementation is effective in normalising plasma zinc levels, thymulin and TSH levels in children with Down syndrome with zinc deficiency (Licastro et al., 1994; Licastro et al., 1992). Clinical evaluation of children with Down syndrome has also shown that zinc supplementation decreased the incidence of infectious diseases, and improved school performance (Franceschi et al., 1988; Licastro et al., 1992).

Hair was used as a biopsy material reflecting the elemental status of the body (Schlegel-Zawadzka, Zachwieja, Huzior-Bajewicz & Pietrzyk, 2002; Misra, Srivastava & Chawla, 1989; Chen, Lin, Lin & Cheng, 1988; Cavadar, Bahceci, Akar, Dincer & Erten, 1991). Trace elements in hair, in particular zinc, are being widely investigated (Deeming & Weber, 1977; Deeming & Weber, 1978; Delves, 1985; Kozielec, Starobrat & Korkowiak, 1994; Litzman, Dastych & Hegar, 1995). Low hair zinc level has been shown to be a good indicator of mild to moderate zinc deficiency (Prasad, 1983; Davies, 1984; Passwater & Cranton, 1983; Hambidge, 1982). Nevertheless hair zinc level is elevated due to impaired hair growth in severe zinc deficiencies (Davies, 1984; Passwater & Cranton, 1983).

In this study we aimed to evaluate the zinc level in children with Down syndrome using hair, which is easily and non-invasively obtained. We also compared the hair zinc levels of children with Down syndrome with typically developing children.
Methods and Materials

Hair samples were collected from the suboccipital area (the back of the skull), by cutting with stainless steel scissors, from 19 children with Down syndrome (age range: 2-6 years old) and 11 age matched typically developing children. The proximal ends of the samples of hair were usually within 2 cm of the scalp, approximately 0.3 g, were used for the analysis. In each of these samples the procedures described below were applied.

Acetone-water wash: Hair samples were placed in 100 ml beakers, covered with bidistilled-deionized water and agitated for 10 minutes with a mechanical shaker. The water was then decanted and replaced with acetone (Merck 13).

The samples were washed four times with acetone each being agitated for a period of 10 minutes and subsequently rinsed with water. After being washed the hair samples were dried at 60°C for 16 hours followed by cooling to room temperature and finally reweighed on an analytical balance.

Wet digestion: hair samples were placed in 100 ml wide mouthed pyrex beakers and left to digest in 5 ml reagent grade nitric acid (Merck 925) overnight. The following morning 10 ml of perchloric acid (Merck 519): nitric acid (1/4 by volume) was added. All samples were covered with watch glasses and refluxed at 200°C for about 8 hours until a clear residue of 2-3 ml remained. After cooling for one hour the digested samples were decanted into 25 ml volumetric flasks and brought to volume with added rinsings of bidistilled-deionized water.

Zinc measurements were done by atomic absorption spectrometry (AAS), using a Pye Unicam SP9. The zinc hollow cathode lamp current was 5 mA and the wavelength was 213.9 nm, slit width 0.5 nm. Results were expressed in parts per million (ppm).

Mean ± standard error of mean values of Zn of both groups in hair were calculated for each group. Anova analysis was used to compare both groups. The 95% confidence interval (CI) on the mean was calculated. Chi-square test was used to compare the cases out of 95% CI. The statistical analyses of the results were calculated in SPSS for Windows.

Results

The results of zinc concentrations of both groups and the comparison of hair zinc levels are presented in Table 1. The mean zinc level was 95.18 ± 56.10 ppm (range: 19.60-191.50) in Down syndrome children, and 208.88 ± 152.37 ppm (range: 52.41-558.53) in typically developing children. Zinc concentrations in hair samples of children with Down syndrome were significantly lower than those of the typically developing children (F(1,28)=8.732, p < .05). The 95% confidence interval of the group of typically developing cases was 102.36 ppm. The number of Down syndrome and typically developing cases out of the lower limit of 95% CI (106.52 ppm) is shown in Table 2. There was no significant difference between the number of Down syndrome and typically developing cases below 95% CI (X²=1.824, df=1, ns).

Discussion

In the present study we used hair to evaluate the zinc levels in children with Down syndrome. We found hair zinc levels of children with Down syndrome were significantly lower than those of the typically developing children. Zinc is an important trace element in metabolism, growth and development and reproduction. It is a constituent of many enzymes. Zinc also plays important roles in nucleic acid metabolism and protein synthesis as well as membrane structure and function. Its deficiency causes impaired growth, poor appetite and physiological changes. Zinc

Table 1. (a) The levels of hair zinc in Down syndrome and typically developing children (b) The mean, SEM and range of zinc levels in Down syndrome and typically developing children
deficiency is also associated with low levels of antibodies. Sustomo & Strblik (1994) reported a high occurrence rate of complex immune and endocrine disorders with thyroid dysregulation in people with Down syndrome, with zinc deficiency playing a considerable role. There is a susceptibility to respiratory infection in children with Down syndrome, and this is one of the major factors in their early mortality. A number of investigations including zinc status have been performed to prevent these infections (Teksen, Sayli, Aydin, Sayal & Işınler, 1998). It has been found that zinc deficiency plays an important role in immunoglobulin concentrations and thyroid function (Franceschi et al., 1988; Licastro et al., 1994; Licastro et al., 1992; Sustomo & Strblik, 1994). However no significant difference was also found between children with Down syndrome with normal zinc levels and low zinc levels regarding the measures of growth hormone secretion, Ig A and Ig G anti-gliadin antibodies, presence of celiac disease, thyroid function tests, CD4/CD8 ratio and total immunoglobulins in another study (Romano et al., 2002). Hair zinc level has also been evaluated in a range of different clinical situations. Mean hair and serum zinc levels were found to be much lower in Indian childhood cirrhosis than the age-matched healthy controls (Mistra, Srivastava & Chawla, 1989). Chen et al. (1988) found the serum and hair zinc controls in obese patients markedly lower than in non-obese patients. Litzman et al. (1995) found a significant decrease in serum zinc levels in common variable immunodeficiency patients. It was found that zinc therapy not only improves the immune system, but also accelerates growth (Napolitano, et al., 1990). Kozielec et al. (1994) showed that it was necessary to supplement trace elements in children with hyperactivity.

Zinc status can be evaluated by using serum, urine, saliva and hair. Although body fluids and tissues are commonly used methods, no correlation has been found between those specimens (Delves, 1985). Hair may be used as a biopsy material reflecting the level of zinc status of the body. Deeming & Weber (1977) evaluated hair analysis for determination of zinc status using rats and reported that hair zinc analysis could be used to aid diagnosis of a deficiency or evaluate dietary intake. Furthermore once incorporated into hair, zinc is no longer in equilibrium with the body and therefore not susceptible to circadian variation (Coker, Cetiner, Sozmen & Ersoz, 1996; Yenigun, Oksel, Bozdogan & Taneli, 1996; Yenigun, Taneli & Kultursay, 1991). The advantages of hair as a source are that it is easy to obtain and stable in storage. It is not affected from the daily variations of food intake. On the other hand, there is a lack of concordance in the results of different laboratories in the assessment and analysis of zinc deficiency (Capel, Spencer, Davies & Levitt, 1985; Deeming & Weber, 1978; Delves, 1985; Lockitch et al., 1989). It has been reported that hair zinc level reflects the body zinc level (Bilir, Kayakirilmaz, Guven, Atik & Ugurlu, 1987; Chen et al., 1985; Kleven, 1970). Deeming & Weber (1978) reported that mineral concentration of hair, serum and diet do not correlate well. Age, sex and body mass index have been reported to be some of the influencing factors that affect the concentration of zinc in the hair (Chen et al., 1985; de Mateo, Perez & Mijan de la Torre, 2000). Kleven (1970) suggested the use of hair as a biopsy material in the evaluation of zinc status by comparing the levels to an age-matched typically developing control group. Our control group was age-matched with the study group, which ranged between 2-6 years old. However, food habits and frequency of intake of different products also influence zinc concentration in hair (Schlegel-Zawadzka et al., 2002; Deeming & Weber, 1978).

These findings indicate that zinc level is influenced by a number of external and internal factors and correction of the zinc level is necessary for the control of biological processes in children with Down syndrome who are vulnerable to zinc deficiency. Therefore zinc level for an individual with Down syndrome may be judged to be deficient after comparing with the age-matched control group from the similar environment. Hair zinc level was investigated in a few studies and ranged between 118 and 152 microgram/g. The mean level of typically developing children was found to be 208.88 ± 152.37 ppm in the present study. Coker et al. (1996) analyzed the hair zinc levels in 24 typically developing children and found a similar mean value to the present study (238.5 microgram/g). The mean hair zinc level in our study was 95.18 ± 56.10 ppm in children with Down syndrome. Accepting the hair zinc values outside of a 95% confidence interval from typically developing children to be abnormal, there is no significant difference between the two groups in the current study. De Mateo et al. (2000) analyzed the zinc status in a healthy, adult, Spanish population. A predictive model of multiple regression was obtained for zinc in hair which is associated with age, sex and BMI. Similar investigations with larger study groups for different populations may be helpful to standardize the hair zinc level, which is an easily obtained and non-invasive biopsy material.

In conclusion, we aimed to investigate the levels of hair zinc in children with Down syndrome by comparison with typically developing controls, therefore we cannot comment on the usefulness of zinc supplementation for children with Down syndrome. Nevertheless zinc supplementation may be useful at least in individuals with Down syndrome with lower hair zinc levels, which is recommended by many authors to reduce the incidence of problems such as infections and endocrinological but further clinical research is required to support this suggestion.

Table 2. The number of cases below / within 95% confidence interval for zinc levels

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<tr>
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<th>Down syndrome</th>
<th>Typically developing</th>
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<tbody>
<tr>
<td>Lower *</td>
<td>10</td>
<td>3</td>
</tr>
<tr>
<td>Within</td>
<td>9</td>
<td>8</td>
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* The number of children below 95% CI

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References


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